FDA 510(k) Premarket Notification

K960475 MODIFIED LUXAR LX-20 FAMILY CO2 LASER SYSTEM

CLEARED

Apr 26, 1996
85 days to decision

K960475 · Product code: **GEX** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k960475/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Powered Laser Surgical Instrument (GEX)
Date received	Feb 1, 1996
Decision date	Apr 26, 1996
Days to decision	85 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Luxar Corp.**
Location	Bothell, WA, US
Contact	JONATHAN S KAHN
510(k) history	17 submissions · 17 cleared · 1988-1996